U.S. Patient and Trademark Office, U.S. DEPARTMENT OF COMMERCE.
Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid CMB control number.

INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
(Not for submission under 37 CFR 1.99)

 Application Number
 10556009

 Filing Date
 2005-11-08

 First Named Inventor
 Johannes P. M. ANSEMS, ET AL.

 Art Unit
 2879

 Examiner Name
 Altoney Docket Number

 NL030899

					U.S.	PATENTS			Remove		
Examiner Initial*	Cite No			Issue [Date Name of Patentee or Applicant of cited Document		Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear				
	1	6169367	B1	2001-01	1-02	MUTO MASA	AKI, ET AL JP				
If you wisl	h to a	i dd additional U.S. Pate	nt citatio	n inform	ation pl	l lease click the	Add button.	_	Add		
			U.S.P	ATENT	APPLI	CATION PUB	LICATIONS		Remove		
Examiner Initial*	Cite No	Publication Number	Kind Code ¹	Publication Date		Name of Pate of cited Docu	entee or Applicant Rele		es,Columns,Lines where evant Passages or Relevant ures Appear		
	1								1		
If you wis	h to a	dd additional U.S. Publ		,		n information p		d button	Remove		_
Examiner Initial*	Cite No	Foreign Document Number ³	Country Code ² i		Kind Code ⁴	Publication	Name of Patente Applicant of cited Document	e or	vhere Rele	or Relevant	T5
	1	0067294	wo		A	2000-11-09	KPENV				
	2	1314317	GB		A	1973-04-18	MATSUSHITA ELE	с			
	3	02168551	JP		A	1990-06-28	SEIKO EPSON CO)RP			_

If you wish	h to a	dd add	dditional Foreign Patent Document citation information please click the Add button Add	
			NON-PATENT LITERATURE DOCUMENTS Remove	
Examiner Cite Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the Initials* No No publisher, city and/or country where publisher.				
	1	ISR,	, WRITTEN OPINION OF THE INTERNATIONAL SEARCHING AUTHORITY PCT//B2/004/05/0606	
If you wis	h to a	dd add	dditional non-patent literature document citation information please click the Add button Add]
			EXAMINER SIGNATURE	
Examiner	Signa	ture	Date Considered	
*EXAMIN	ER: Ir	itial if	if reference considered, whether or not citation is in conformance with MPEP 609. Draw line th	rough a

1 See Kind Codes of USPTO Patent Documents at wear USPTO_GOLY or MEPP 901.64. If Enter office that issued the document, by the low-letter of Kind Standard \$7.33, "3 or duptamese patent outcoment, the robusto of the year of the inspect may precede the serial runber of the opatent document, "4 Kind of document by the appropriate symbols as indicated on the document under WIPO Standard \$7.16 if possible. "Applicant is to place a check mark here if Enginh Imaguage the resistation is attacked."

citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		10556009
Filing Date		2005-11-08
First Named Inventor Johan		ines P. M. ANSEMS, ET AL
Art Unit		2879
Examiner Name		
Attorney Docket Number		NL030699

CERTIFICATION STATEMENT

Please see 37	CFR 1.97 and	1.98 to make the a	ppropriate selection(s):
---------------	--------------	--------------------	--------------------------

That each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filling of the information disclosure statement. See 37 CFR 1.97(e/11).

OR

That no item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification after making reasonable inquity, no item of information contained in the information disclosure statement was known to any individual designated in 37 CFR 156(c) more than three months prior to the filing of the information disclosure statement. See 37 CFR 157(e).

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.
- ▼ None

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

_			
Signature	/Frank J. Keegan/	Date (YYYY-MM-DD)	2007-08-31
Name/Print	Frank J. Keegan	Registration Number	50,145

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is for life railed by the USPTO to process) an application. Confidentiality is governed by \$5 U.S. C. 12.0 and 3T CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case: Any comments on the amount of time you require to complete his form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. operatment of Commence, P.O. 8bx 1449, Alexandriv, V.S. 2311-1450, D.O. NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. 8bx 1459, Alexandriva, V.S. 2311-1450.

Privacy Act Statement

The Privacy Act of 1974 (P. L. 93-579) requires that you be given certain information in connection with your submission of the stackhold from related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, places be advised that (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) familishing of the information solicided is couldrain; and (3) the primoral pursuance for which the information is used by the U.S. Patient and Trademan Coffice is to process and/or examine your submission related to a patient agricultant or patient. If you do not furnish the requested process and/or examine your submission related to a patient agricultant or patient. If you do not furnish the requested results of the patient of the patient and the patient of the patient

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement negotiation.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record pertains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974, as amended, pursuant to 5 U.S.C. 552(m).
 - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
 - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uturing an insection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 4d U.S.C. 2904 and 2905. Such disclosure shall be made in accordance with the GSA requisions governing inseption of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the
 application pursuant to 35 U.S.C. 12(2) to rissuance of a patent pursuant to 35 U.S.C. 151. Further, a record may be
 disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record was filled in application
 which became abandoned or in which the proceedings were terminated and which application is referenced by either a
 published application, an application open to public inspections or as issued patent.
 - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.